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NOVEL COPOLYMER, PHOTORESIST COMPOSITION, AND PROCESS FOR
FORMING RESIST PATTERN WITH HIGH ASPECT RATIO

BACKGROUND OF THE INVENTION

Field of the Invention

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The present invention relates to a chemically amplified photoresist composition that is excellent in transparency to light with wavelengths in the deep ultraviolet region (deep UV light) such as KrF or ArF laser light, and exhibits high sensitivity and definition, to a novel copolymer that is suitable for the preparation of the photoresist composition, and to a process for the formation of a resist pattern with a high aspect ratio. Specifically, the present invention relates to a chemically amplified photoresist composition that is suitable for a process for the formation of a resist pattern with a high aspect ratio using a silylation agent, and to a process for the formation of a resist pattern with a high aspect ratio.

Description of the Related Art

To form an ultrafine resist pattern of not more than 0.35 μm , specifically of not more than 0.25 μm , a variety of highly sensitive photoresist compositions having satisfactory transparency to deep UV light such as KrF or

ArF laser light have been reported.

Among them, "chemically amplified" photoresist compositions containing a substance carrying an acid-decomposable group, and a photosensitive acid generator are predominant. However, demands have been made to further improve the sensitivity and definition of these chemically amplified photoresist compositions.

Additionally, the use of this type of photoresist compositions alone cannot significantly form a resist pattern with a good shape, especially a hole-type resist pattern with a high aspect ratio and a good shape. As a possible solution to this problem, a technique has been reported in which a second film or coating composed of a silylation agent is formed on the surface of a resist pattern, and using this second film, a resist pattern with a high aspect ratio is formed.

This technique using a silylation agent is reported, for example, in Microelectronic Engineering 11 (1990) 531-534 (Reference 1), and is performed substantially in the following manner.

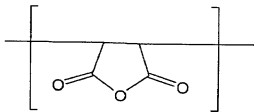
Initially, a resist film (bottom resist) is formed on a substrate, and this bottom resist is resistant to etching treatment for the substrate. Another resist film is then formed on the bottom resist, using a photoresist composition

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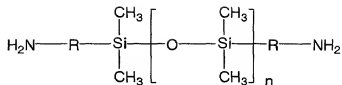
containing, for example, a polymer having a repeating unit represented by Formula (V) below, and the formed resist film is selectively patterned by exposure and developing to thereby form a resist pattern.

Next, a solution of, for example, a compound represented by Formula (VIII) below (silylation agent) is applied onto the resist pattern and is then rinsed to thereby form a silylation coating on the resist pattern, and the resulting silylation coating is resistant to corrosion induced by oxygen plasma etching.

The underlayer bottom resist is removed by etching using, as a mask, the resist pattern carrying the silylation coating to thereby form a resist pattern with a high aspect ratio.



(V)



(VIII)

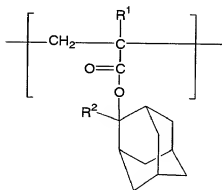
Such photoresist compositions that can be applied to this process comprising the above operations must essentially react with a silylation agent to form a silylation coating and must have high sensitivity and definition.

Japanese Patent Laid-Open No. 5-11450 (Reference 2) discloses, as the aforementioned photoresist compositions, a photoresist composition containing a polymer and a photosensitive acid generator, which polymer comprises a (meth)acrylic tert-butyl ester group and a maleic anhydride functional group, and a photoresist composition containing a polymer and a photosensitive acid generator, which polymer comprises a (meth)acrylic tert-butyl ester group, a maleic anhydride functional group, and an allyltrimethylsilane group.

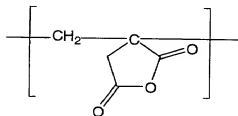
These compositions have satisfactory transparency to deep UV light and have high sensitivity, but their definition must be further improved.

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Japanese Patent Laid-Open No. 11-212265 (Reference 3) discloses a photoresist composition comprising a polymer and a photosensitive acid generator, which polymer has a repeating unit represented by Formula (III) and a repeating unit represented by Formula (VII):



(III)



(VII)

wherein R^1 is a hydrogen atom or a methyl group; and R^2 is an alkyl group having from 1 to 4 carbon atoms.

Even this composition must be further improved in sensitivity and definition.

Additionally, when this composition is applied to the process using a silylation agent, it cannot significantly react with the silylation agent to thereby fail to form a silylation coating, and a hole pattern with a high aspect ratio cannot be significantly obtained.

SUMMARY OF THE INVENTION

Accordingly, an object of the present invention is to provide a chemically amplified photoresist composition that has satisfactory transparency, high sensitivity and definition in the field of photolithography using a deep UV light source such as KrF or ArF laser, and to provide a novel copolymer that is suitable for the preparation of the photoresist composition.

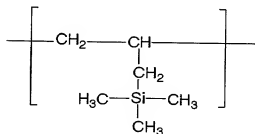
Another object of the present invention is to provide a chemically amplified photoresist composition that is suitable for a process for the formation of a resist pattern with a high aspect ratio using a silylation agent, and to provide a process for the formation of a resist pattern with a high aspect ratio.

After intensive investigations, the present inventors have found that the above objects can be achieved by the use of novel copolymers having specific repeating units.

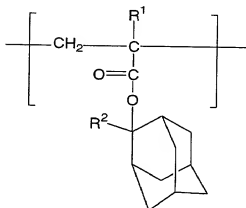
Specifically, the present invention provides, in an

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aspect, a novel copolymer including a repeating unit (B) derived from an unsaturated carboxylic anhydride, a repeating unit (C) represented by Formula (II), and a repeating unit (D) represented by Formula (III):



(II)



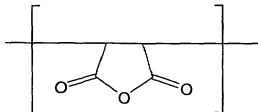
(III)

wherein R¹ is a hydrogen atom or a methyl group; and R² is an alkyl group having from 1 to 4 carbon atoms.

In the novel copolymer, repeating unit (B) is preferably a repeating unit (B-1) derived from an

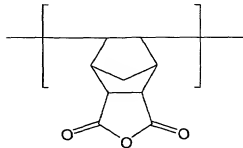
unsaturated cyclic carboxylic anhydride.

Repeating unit (B-1) is preferably a unit (B-2) represented by Formula (V):



(V)

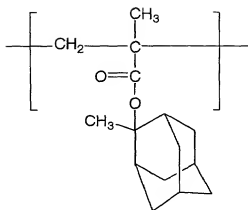
Alternatively, repeating unit (B-1) is preferably a unit (B-3) represented by Formula (XVI):



(XVI)

In the novel copolymer, repeating unit (D) may be a unit (D-1) represented by Formula (VI):

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(VI)

The content of repeating unit (B) is preferably equal to or more than 15% and equal to or less than 60% of all repeating units constituting the novel copolymer.

The content of repeating unit (C) is preferably equal to or more than 10% and equal to or less than 40% of all repeating units constituting the novel copolymer.

The content of repeating unit (D) is preferably more than 0% and equal to or less than 40% of all repeating unit constituting the novel copolymer.

The novel copolymer may have a weight average molecular weight (Mw) in terms of polystyrene of from 7000 to 30000 and a molecular-weight distribution (Mw/Mn, where Mn is a

number average molecular weight) of equal to or less than 3.5.

In another aspect, the present invention provides a photoresist composition including the novel copolymer, a photosensitive acid generator, and an organic solvent.

In the photoresist composition, the photosensitive acid generator is preferably a triphenylsulfonium-based onium salt.

The organic solvent is preferably propylene glycol monomethyl ether acetate (PGMEA).

In addition and advantageously, the present invention provides a process for forming a resist pattern with a high aspect ratio, which process includes the steps of (a) applying a first resist on a substrate and drying the applied first resist to thereby form a first resist layer, applying the photoresist composition the first resist layer and drying the applied photoresist composition to thereby form a second resist layer; (b) exposing the second resist layer to imaging radiation, subjecting the exposed second resist layer to a heat treatment, and dissolving and removing exposed portions or unexposed portions of the second resist layer by developing in an alkaline aqueous solution to thereby form a resist pattern; (c) applying a silylation agent onto the formed resist pattern, rinsing the

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applied resist pattern to thereby enlarge the resist pattern and to form a silylation coating on the resist pattern, which silylation coating is resistant to corrosion induced by oxygen-containing plasma etching; and (d) etching the first resist layer under the second resist layer with oxygen-containing plasma by using, as a mask, the enlarged resist pattern carrying the silylation coating.

DETAILED DESCRIPTION OF THE INVENTION

[Novel Copolymers]

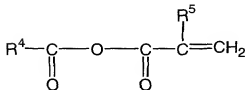
Novel copolymers of the present invention can be synthetically obtained by a known polymerization reaction using specific comonomers. Repeating units (B), (C) and (D) (hereinafter briefly referred to as "Ingredients (B), (C) and (D)") will be illustrated in further detail below, as constitutional comonomers for the preparation of the novel copolymers.

[Ingredient (B)]

Ingredient (B) constituting the invented copolymers is a repeating unit derived from an unsaturated carboxylic anhydride and is an essential unit, since the repeating unit serves to react with a silylation agent (e.g., a siloxane polymer having an amino group) to thereby form a silylation coating in the process using the silylation agent.

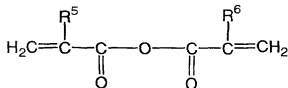
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Comonomers corresponding to Ingredient (B) are not specifically limited and include, for example, unsaturated carboxylic anhydrides described in Japanese Patent Laid-Open Nos. 2-282746, 2-308255, 2-308256, 5-9231, 5-11450, 5-11456, and 11-212265. Such unsaturated carboxylic anhydrides include, for example, compounds represented by Formulae (IX) and (X), and other unsaturated acyclic carboxylic anhydrides; and compounds represented by Formulae (XI), (XII), (XIII), (XIV), and (XV), and other unsaturated cyclic carboxylic anhydrides:



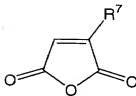
(IX)

wherein R⁴ is an alkyl group having from 1 to 5 carbon atoms or an aryl group; and R⁵ is a hydrogen atom or an alkyl group having from 1 to 4 carbon atoms,



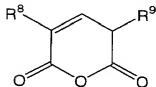
(X)

wherein each of R^5 and R^6 is independently a hydrogen atom or an alkyl group having from 1 to 4 carbon atoms,



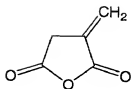
(XI)

wherein R^7 is a hydrogen atom or an alkyl group having from 1 to 4 carbon atoms,

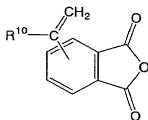


(XII)

wherein each of R^8 and R^9 is independently a hydrogen atom or an alkyl group having from 1 to 4 carbon atoms,

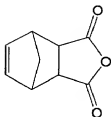


(XIII)



(XIV)

wherein R^{10} is a hydrogen atom or an alkyl group having from 1 to 4 carbon atoms.



(XV)

Among these comonomers, maleic anhydride is typically preferable for its high reactivity with a silylation agent and satisfactory transparency to deep UV light such as KrF or ArF laser light. Preferred Ingredients (B) for use in

the invented copolymers are repeating units (B-2) and (B-3) represented by Formulae (XI) and (XV) which are derived from unsaturated cyclic carboxylic anhydrides.

[Ingredient (C)]

Ingredient (C) constituting the invented copolymers is a repeating unit represented by Formula (II) and serves to enhance reactivity with a silylation agent (e.g., a siloxane polymer having an amino group) in the process using the silylation agent. Additionally, the resulting photoresist composition obtained by using a copolymer containing Ingredient (C) has improved sensitivity and definition.

A comonomer corresponding to Ingredient (C) is allyltrimethylsilane.

[Ingredient (D)]

Ingredient (D) constituting the invented copolymers is a repeating unit represented by Formula (III), which has an 2-alkyl-2-adamantyl group as an acid-decomposable group and is satisfactory in transparency to deep UV light such as KrF or ArF laser light. The substituent R^1 in the formula is a hydrogen atom or a methyl group, and R^2 is an alkyl group having from 1 to 4 carbon atoms. The resulting photoresist composition prepared by using a copolymer containing Ingredient (D) has further improved sensitivity and definition.

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Comonomers corresponding to Ingredient (D) include, for example, 2-methyl-2-adamantyl acrylate, 2-ethyl-2-adamantyl acrylate, 2-propyl-2-adamantyl acrylate, 2-butyl-2-adamantyl acrylate, 2-methyl-2-adamantyl methacrylate, 2-ethyl-2-adamantyl methacrylate, 2-propyl-2-adamantyl methacrylate, and 2-butyl-2-adamantyl methacrylate. Among them, 2-methyl-2-adamantyl methacrylate is typically preferred to yield higher sensitivity and definition of the resulting photoresist composition. These comonomers corresponding to Ingredient (D) can be synthetically obtained by known processes as described in, for example, Japanese Patent Laid-Open No. 11-212265. For example, these comonomers can be obtained by an esterification reaction of an 2-alkyl-2-adamantanol with acryloyl chloride or methacryloyl chloride.

[Contents of Ingredients (B), (C) and (D)]

The content of Ingredient (B) preferably falls within a range from 15% to 60% inclusive, and more preferably from 35% to 50% inclusive, of all the repeating units constituting the novel copolymer.

If the content of Ingredient (B) is less than 15%, the reactivity with the silylation agent may be insufficient, and if it exceeds 60%, Ingredient (B) may be readily converted into a dicarboxylic acid moiety by a reaction with moisture in the air or in the photoresist film to thereby

polymerization reaction with a polymerization initiator.

Such polymerization initiators include, but are not limited to, those described in, for example, Japanese Patent Laid-Opens No. 5-11450 and No. 11-212265, such as azobisisobutyronitrile and dimethyl-2,2-azoisobisbutyrate. Among them, azobisisobutyronitrile is typically preferred, since this compound can enhance the polymerization reaction to thereby yield a copolymer having narrow molecular-weight distribution.

An example of the production process of the invented novel copolymers will be schematically illustrated below, but the invention is not limited thereto.

(Step 1)

Initially, the material comonomers, polymerization initiator, and additives are added to an organic solvent (e.g., ethyl acetate) in a reactor and are dissolved in the organic solvent by stirring at room temperature for several ten minutes.

(Step 2)

The resulting mixture is heated to a temperature from about 60°C to about 75°C, and is stirred at this temperature for several ten hours.

(Step 3)

After the completion of stirring, heating procedure is

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copolymer, a photosensitive acid generator, and an organic solvent. The content of the novel copolymer is preferably from 1 to 30% by weight and more preferably from 3 to 10% by weight relative to the total weight of the composition.

[Photosensitive Acid Generator]

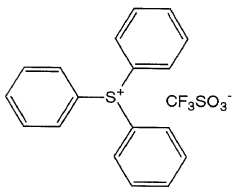
When the invented photoresist composition is used for the formation of a resist pattern, the photosensitive acid generator liberates an acid in exposed portions in the step of exposing a photoresist film to imaging radiation. Additionally, a subsequent heat treatment permits an "acid-decomposable group" (e.g., an 2-alkyl-2-adamantyl group) in the copolymer skeleton of the composition to eliminate, which acid-decomposable group is decomposed by the catalytic reaction of the acid and becomes soluble in alkali. Consequently, exposed portions of the resist film are dissolved in a developer solution, an alkaline aqueous solution, to thereby yield a resist pattern.

Photosensitive acid generators for use in the present invention include, but are not specifically limited to, those described in, for example, Japanese Patent Laid-Open Nos. 5-11450 and 11-212265, such as onium salt compounds, organohalogen compounds, sulfone compounds, and sulfonate compounds. Among them, onium salt-based photosensitive acid generators are preferred, since the resulting photoresist

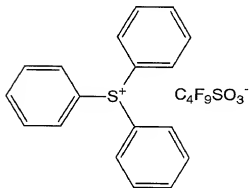
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compositions have high sensitivity and can form a resist pattern with a good shape, of which triphenylsulfonium-based onium salts are typically preferred for high sensitivity.

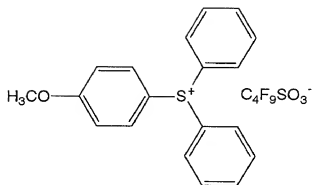
Such triphenylsulfonium-based onium salts include, for example, the following compounds represented by Formulae (E-1) to (E-3):



(E-1)



(E-2)



(E-3)

The amount of the photosensitive acid generator is preferably from 0.01 to 3.0% by weight and more preferably from 0.03 to 0.8% by weight relative to the total weight of the composition.

[Organic Solvent]

Organic solvents for use in the invention include, but are not specifically limited to, acetone, methyl ethyl ketone, cyclohexanone, methyl isoamyl ketone, 2-heptanone, and other ketones; ethylene glycol, propylene glycol, diethylene glycol, ethylene glycol monoacetate, propylene glycol monoacetate, diethylene glycol monoacetate, or monomethyl ethers, monoethyl ethers, monopropyl ethers, monobutyl ethers, or monophenyl ethers of these compounds, and other polyhydric alcohols and derivatives thereof; dioxane and other cyclic ethers; and ethyl lactate, methyl acetate, ethyl acetate, butyl acetate, methyl pyruvate,

ethyl pyruvate, methyl methoxypropionate, ethyl ethoxypropionate, and other esters. Each of these organic solvents can be used alone or in combination.

Among them, propylene glycol monomethyl ether acetate (PGMEA) is preferred, since this solvent has good coating performance, low hygroscopicity, and a high boiling point. Solvents having high hygroscopicity are not desirable, since the resulting photoresist composition takes in moisture in the air to thereby deteriorate the copolymer. Additionally, solvents having a low boiling point are also undesirable, since the amount of solvent remained in the formed photoresist film is low and the photosensitive acid generator does not efficiently diffuse in the procedure of, for example, post-exposure baking (PEB).

[Other Additives]

The invented photoresist compositions may further comprise known or conventional additives according to necessity, within ranges not adversely affecting the objects of the present invention. Such additives include, for example, dissolution inhibitors, other resins, stabilizers, organic polymeric compounds, thermopolymerization inhibitors, sensitizers, improvers for adhesion to substrates, surfactants, and dyes.

The invented photoresist compositions are suitable as

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positive photoresist compositions, but can be used as negative photoresist compositions for the formation of negative resist patterns, by incorporating acid-crosslinkable materials (substances which are crosslinked by action of an acid generated from a photosensitive acid generator) into the photoresist compositions.

The invented composition is preferably used, for example, in the following manner. Initially, a solution of the (positive) photoresist composition is applied on a substrate such as a silicon wafer, and is then dried to thereby form a photosensitive layer; and the photosensitive layer is exposed to light through a patterned photomask.

Next, the exposed portions of the photosensitive layer are subjected to post-exposure baking (PEB) and are then dissolved and removed by dipping the substrate in a developer solution, for example, an alkaline aqueous solution such as a 1% to 10% by weight tetramethylammonium hydroxide (TMAH) aqueous solution, thus forming an image (resist pattern) being in exact accordance with the mask pattern. To further improve the definition of the resist pattern, an anti-reflection coating is preferably formed between the substrate and the photosensitive layer composed of the invented composition.

[Process for Forming Resist Pattern with High Aspect Ratio]

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The invented photoresist compositions are specifically suitable for the process for the formation of a resist pattern using a silylation agent. In the process, adjacent resist patterns (line width) are enlarged and grooves (space width) are narrowed by using a silylation agent to thereby form a resist pattern with a high aspect ratio, and the invented photoresist compositions can be advantageously used in this process. The process will be schematically illustrated below, but the invention is not limited thereto.

(Step a)

A first resist that can be smoothed is applied on a substrate and is dried to thereby form a first resist layer (herein after referred to as "bottom resist", which will be described later), and the invented photoresist composition that can form a resist pattern by photolithography is applied onto the first resist layer and is dried to thereby form a second resist layer (hereinafter referred to as "top resist").

(Step b)

Subsequently, the top resist is exposed to imaging radiation and is then subjected to a heat treatment, and the exposed portions (in case of a positive photoresist composition) or unexposed portions (in case of a negative photoresist composition) are dissolved and removed by

suitable. By heating the applied bottom resist on the substrate, cross-linking of the novolak resin occurs to thereby improve resistance to etching and to inhibit dissolution of the bottom resist and the top resist (the invented photoresist composition) in each other.

[Silylation Agent]

Silylation agents for use in the above process are enlarging reagents that can be bonded with the resist pattern of the top resist (the invented photoresist composition) and can increase the volume of the resist pattern. Additionally, the resist pattern carrying a silylation coating composed of the silylation agent can become a layer that is very resistant to corrosion induced by dry developing in oxygen-containing plasma etching for etching the bottom resist.

Silylation agents having a Si-O (siloxane) structure are preferred as the silylation agents, since a nonvolatile oxide is formed from a silicon-containing group in the silylation coating during oxygen-containing plasma etching to thereby enhance corrosion resistance, and the siloxane compounds are highly thermally stable to thereby avoid pattern deformation during the etching step at high temperatures. These siloxane compounds can be easily directly converted into silicon dioxide and produce little

volatile organosilicon compound in oxygen-containing plasma. Among them, siloxane compounds having an amino group are preferred, such as aminosiloxane, diaminosiloxane, and bisaminopropylpolydimethylsiloxane. The amino group in the silylation agent rapidly reacts with Ingredient (B) of the copolymer in the invented photoresist composition on the surface of the resist pattern of the top resist to increase the volume of the resist pattern to thereby narrow the space pattern.

Commercially available agents can be used as the silylation agents. For example, a preferred silylation agent solution for use in the invention is a 1% by weight bisaminopropylpolydimethylsiloxane solution prepared by dissolving bisaminopropylpolydimethylsiloxane (available from Shin-Etsu Chemical Co., Ltd., under the trade name of "X-22-161 AS") in an aqueous 1-hexanol solution.

BRIEF DESCRIPTION OF THE DRAWINGS

Fig. 1 is a diagram showing the spectrum of Copolymer 1 obtained in Synthesis Example 1, measured by Fourier transform infrared spectrophotometry (FT-IR); and

Fig. 2 is a diagram showing the chart of Copolymer 1 obtained in Synthesis Example 1, measured by gel permeation chromatography (GPC).

EXAMPLES

The present invention will be illustrated in further detail with reference to several examples and comparative examples below, which are not intended to limit the scope of the invention.

The physical properties of the photoresist compositions were determined according to the following methods.

(1) Sensitivity Evaluation

Initially, a novolak resin-quinonediazide positive photoresist composition (available from Tokyo Ohka Kogyo Co., Ltd., under the trade name of "THMR-iP5700) was applied on a silicon wafer, and was cured by heat to thereby form a resist film (bottom resist) 0.6 μm thick. This was used as a substrate in the subsequent procedure. A sample was applied onto the substrate using a spinner, and was dried on a hot plate at 110°C for 90 sec. to form a resist film 0.2 μm thick. The resist film was then irradiated through a mask corresponding to a hole diameter of 0.23 μm and a duty ratio of 1:0.57 (interval between holes: 0.13 μm) using a reducing-type projection aligner NSR-S203B (available from Nikon Corporation, Japan; NA=0.60, δ =0.68) with increasing bias from 10 mJ/cm² at intervals of 1 mJ/cm². The film was then post-exposure baked (PEB) at 130°C for 90 sec.; was subjected to developing in a 2.38% by weight

tetramethylammonium hydroxide aqueous solution at 23°C for 40 sec., was washed with water for 15 sec., and was dried. In this procedure, the sensitivity was defined as the exposure time period (Eop) (mJ/cm^2) to reproduce a 0.18- μm hole pattern.

(2) Definition

The definition was defined as the minimum mask size (μm) in the critical definition to separate the bottom of the pattern at an exposure of Eop in the procedure of Evaluation (1).

(3) Focal Depth Range Properties

A sample was subjected to exposure and developing in the same manner as in Evaluation (1), except that the focus was shifted up and down at an exposure of Eop as the standard exposure. The resulting hole pattern was subjected to a scanning electron micrographic (SEM) observation. Based upon the SEM photograph, the focal depth range property was defined as the maximum value (μm) of the focal shift (defocus) to reproduce a 0.18- μm hole pattern when the hole pattern was formed under bias using a 0.23- μm mask.

SYNTHESIS EXAMPLE 1: Synthesis of Copolymer 1

To 69.4 parts by weight of ethyl acetate in a reactor, 15.9 parts by weight (0.36 mole) of 2-methyl-2-adamantyl methacrylate, 9.9 parts by weight (0.54 mole) of maleic

anhydride, 6.4 parts by weight (0.30 mole) of allyltrimethylsilane, and 0.6 part by weight of azobisisobutyronitrile were added, and were stirred at room temperature (25°C) for 60 minutes.

Next, the resulting mixture was heated to about 70°C and was stirred at this temperature for about 22 hours.

To the resulting reaction mixture, 11.5 g (0.36 mole) of a 2.0% by weight methanol aqueous solution was added dropwise, and the mixture was further stirred for about 22 hours.

After the completion of stirring, the heating operation was stopped, and the reaction mixture was cooled to about 30°C and was then added dropwise to petroleum benzine with stirring to thereby precipitate a copolymer.

The precipitated copolymer was rinsed several portions of petroleum benzine and was dried in a vacuum dessicator set at about 60°C for 3 hours to thereby yield Copolymer 1.

Copolymer 1 was subjected to FT-IR assay by the KBr method using an FT-IR system (available from The Perkin-Elmer Corporation, under the trade name of "SPECTRUM 2000").

The results are shown in Fig. 1.

Fig. 1 shows that the characteristic band of adamantyl moiety of 2-methyl-2-adamantyl methacrylate in the vicinity of 1100 cm^{-1} , the absorption of C=O stretching vibration of

maleic anhydride in the vicinity of 1780 cm^{-1} , and the absorption of the deformation vibration of Si-CH_3 of allyltrimethylsilane in the vicinity of 1250 cm^{-1} were respectively observed.

Separately, Copolymer 1 was subjected to gel permeation chromatographic (GPC) analysis by the R.I. method using a GPC system (available from Shodex, under the trade name of "GPC-SYSTEM-11").

The results are shown in Fig. 2.

Fig. 2 shows that Copolymer 1 was a polymer having a weight average molecular weight (M_w) in terms of polystyrene of about 18700 and a molecular-weight distribution (M_w/M_n) of about 1.67.

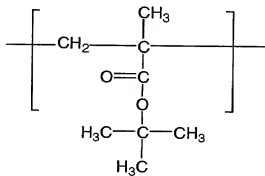
SYNTHESIS EXAMPLES 2 TO 4: Synthesis of Copolymers 2 to 4

Copolymers 2 to 4 containing the repeating units indicated in Table 1 were synthetically obtained in the same manner as in Synthesis Example 1, except that the types and compositional ratios of individual comonomers were varied.

COMPARATIVE SYNTHESIS EXAMPLES 1 AND 2: Synthesis of Copolymers 5 and 6

Copolymers 5 and 6 containing the repeating units indicated in Table 1 were synthetically obtained in the same manner as in Synthesis Example 1, except that the types and

compositional ratios of individual comonomers were varied. Copolymer 5 as a comparative example was obtained by using tert-butyl methacrylate as a comonomer. The repeating unit derived from tert-butyl methacrylate is a unit represented by Formula (X-1) below.



(X-1)

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Table 1

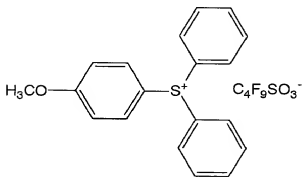
Example No. (Copolymer) (Mw) (Mw/Mn)	(B) (mole %)	(C) (mole %)	(D) (mole %)	(X) (mole %)
Synthesis Example 1 (Copolymer 1) (18700) (1.67)	B-2 (45)	C (25)	D-1 (30)	none
Synthesis Example 2 (Copolymer 2) (10000) (1.6)	B-3 (45)	C (25)	D-1 (30)	none
Synthesis Example 3 (Copolymer 3) (15000) (1.9)	B-2 (45)	C (20)	D-1 (35)	none
Synthesis Example 4 (Copolymer 4) (16800) (1.9)	B-2 (45)	C (30)	D-1 (25)	none
Comp. Syn. Ex. 1 (Copolymer 5) (20000) (2.0)	B-2 (50)	C (27.5)	none	X-1 (22.5)
Comp. Syn. Ex. 2 (Copolymer 6) (20000) (2.0)	B-2 (70)	none	D-1 (30)	none

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EXAMPLES 1 TO 4 AND COMPARATIVE EXAMPLES 1 AND 2

Each of Copolymers 1 to 6 synthetically obtained in Synthesis Examples 1 to 4 and Comparative Synthesis Examples 1 and 2 was dissolved in PGMEA to thereby yield a 7% by weight polymer solution.

In 30 g of the above-prepared polymer solution, 0.158 g of a triphenylsulfonium-based onium salt represented by Formula (E-3) below was dissolved, and the resulting solution was filtrated though a membrane filter of 0.1 to 0.2 μm pore size to thereby yield a series of photoresist compositions.



(E-3)

Evaluations (1) to (3) were conducted on the photoresist compositions prepared according to Examples 1 to 4 and Comparative Examples 1 and 2, and the results are shown in Table 2.

Table 2

	Sensitivity (mJ/cm ²)	Definition (μm)	Focal depth range property (μm)
Example 1	45	0.16	0.6
Example 2	45	0.17	0.6
Example 3	30	0.16	0.5
Example 4	55	0.17	0.5
Comp. Ex. 1	40	0.18	0.6
Comp. Ex. 2	80	0.18	0.4

SYNTHESIS EXAMPLE 5: Preparation of silylation agent solution

Bisaminopropylpolydimethylsiloxane (available from Shin-Etsu Chemical Co., Ltd., under the trade name of "X-22-161 AS") was dissolved in a 98.5% by weight 1-hexanol aqueous solution to thereby yield a 1% by weight silylation agent solution.

EXAMPLE 5: Formation of a resist pattern with a high aspect ratio using the silylation agent

A hole pattern [a hole pattern 0.18 μm wide] was formed on a substrate at depth of focus of 0 in the same manner as in Evaluation (3) in Example 1, and the above-prepared silylation agent solution was carefully placed as a heap on

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the substrate carrying the hole pattern, and was allowed to stand for 30 seconds, and the substrate was rinsed with isopropanol for 15 seconds.

The cross section (profile) of the resist pattern after the silylation was subjected to SEM photographic observation and was found that the initial hole diameter, $0.18\text{ }\mu\text{m}$, was narrowed to $0.14\text{ }\mu\text{m}$ by the formation of a silylation coating and that the hole profile was not deteriorated.

COMPARATIVE EXAMPLE 3: Formation of a resist pattern with a high aspect ratio using the silylation agent

A hole pattern [a hole pattern $0.18\text{ }\mu\text{m}$ wide] was formed on a substrate at depth of focus of 0 in the same manner as in Evaluation (3) in Comparative Example 1, and the above-prepared silylation agent solution was carefully placed as a heap on the substrate carrying the hole pattern, and was allowed to stand for 30 seconds, and the substrate was rinsed with isopropanol for 15 seconds.

The cross section (profile) of the resist pattern after the silylation was subjected to SEM photographic observation and was found that the initial hole diameter, $0.18\text{ }\mu\text{m}$, was narrowed to $0.15\text{ }\mu\text{m}$ by the formation of the silylation coating but that the hole profile was deteriorated.

COMPARATIVE EXAMPLE 4: Formation of a resist pattern with a high aspect ratio using the silylation agent

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A hole pattern [a hole pattern 0.18 μm wide] was formed on a substrate at depth of focus of 0 in the same manner as in Evaluation (3) in Comparative Example 2, and the above-prepared silylation agent solution was carefully placed as a heap on the substrate carrying the hole pattern, and was allowed to stand for 30 seconds, and the substrate was rinsed with isopropanol for 15 seconds.

The cross section (profile) of the resist pattern after the silylation was subjected to SEM photographic observation and was found that no silylation coating was formed and that the hole diameter was not narrowed.

ADVANTAGES

The present invention can provide a chemically amplified photoresist composition that has satisfactory transparency, high sensitivity and definition in the field of photolithography using a deep UV light source such as KrF or ArF laser, and provide a novel copolymer that is suitable for the preparation of the photoresist composition.

Additionally, the present invention can provide a chemically amplified photoresist composition that is suitable for a process for the formation of a resist pattern with a high aspect ratio using a silylation agent, and provide a process for the formation of a resist pattern with a high aspect ratio.

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Other embodiments and variations will be obvious to those skilled in the art, and this invention is not to be limited to the specific matters stated above.

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